

IN THE UNITED STATES PATENT AND TRADEMARK OFFICE

Applicants: Baker et al. Docket No: 39780-2830P1C8  
Serial No: 10/006,041 Group Art Unit: 1647  
Filed: December 06, 2001 Examiner: Rachel K. Hunnicutt  
For: **SECRETED AND TRANSMEMBRANE POLYPEPTIDES AND NUCLEIC  
ACIDS ENCODING THE SAME**

Commissioner for Patents  
P.O. Box 1450  
Alexandria, Virginia 22313-1450

**DECLARATION OF NAPOLEONE FERRARA, Ph.D.,**  
**AUDREY GODDARD, Ph.D., PAUL J. GODOWSKI, Ph.D.,**  
**AUSTIN GURNEY, Ph.D., JAMES PAN, Ph.D., COLIN K. WATANABE and**  
**WILLIAM I. WOOD, Ph.D. UNDER 37 CFR 1.131**

We, Napoleone Ferrara, Ph.D., Audrey Goddard, Ph.D., Paul J. Godowski, Ph.D., Austin Gurney, Ph.D., James Pan, Ph.D., Colin K. Watanabe and William I. Wood, Ph.D. declare and say as follows:

1. We are the inventors of the above-identified application.
2. We have read and understood the claims pending in this application, and are aware that the claims have been rejected as anticipated by U.S. Patent No. 6,525,174 (Young *et al.*, issue date February 25, 2003 and effective filing date June 4, 1998) and U.S. Patent Publication No. 2003/0096951 (Jacobs *et al.*, publication date May 22, 2003 and effective filing date August 14, 1998).
3. The polypeptide designated as PRO1244 (SEQ ID NO:130) claimed in the above-identified application in the United States was sequenced, cloned and homology to implantation-associated protein identified prior to June 4, 1998.
4. At the time the PRO1244 polypeptide was cloned and sequenced, one of the inventors, Austin Gurney, Ph.D., was responsible for overseeing the cloning of cDNAs which

encoded novel polypeptides, including the cDNA that encoded PRO1244 polypeptide (SEQ ID NO:130) claimed in the above-identified application.

5. At the time the PRO1244 polypeptide was cloned and sequenced, one of the inventors, Audrey Goddard, Ph.D., was, and still is, responsible for overseeing the sequencing of novel polypeptides, including the PRO1244 polypeptide (SEQ ID NO:130) claimed in the above-identified application.
6. At the time the PRO1244 polypeptide was cloned and sequenced, one of the inventors, William I. Wood, was, and still is, responsible for overseeing the homology searches for the novel polypeptides, including that for the PRO1244 polypeptide (SEQ ID NO:130) encoded by the cDNA claimed in the above-identified application.
7. A cDNA clone, referred to as DNA64883-1526 in the above-identified application, was identified as encoding the PRO1244 polypeptide.
8. The full length of the cDNA clone is shown in Figure 73 of the above-identified application. The full-length cDNA sequence has 2213 nucleotide residues. The full length of the PRO1244 peptide encoded by DNA64883-1526 is shown in Figure 74 of the above-identified application. The full-length PRO1244 polypeptide has 335 amino acid residues.
9. Copies of the pages from the GSeqEdit database which report the cloning and sequencing data for the PRO1244 polypeptide sequence and its encoding nucleic acid sequence are attached to this declaration (with the dates redacted) as Exhibit A.
10. The GSeqEdit report shows the full-length nucleic acid sequence for DNA-64883-1526 (identified as "DNA-64883") and the full-length PRO1244 polypeptide encoded by DNA 64883. Both the DNA-64883 and the PRO1244 polypeptide sequences were obtained prior to June 4, 1998.
11. The DNA-64883 sequence shown in the GSeqEdit report is identical to that of SEQ ID NO: 129 disclosed in the above-identified application.
12. The beginning of the cDNA sequence corresponding to SEQ ID NO: 129 in the above-identified application is shown on page 1 of the GSeqEdit database report and

the location of the first nucleotide is marked with "insert starts here" and an arrow. The location of the last nucleotide corresponding to SEQ ID NO: 129 is shown on page 11 and is marked with an arrow.

13. The amino acid sequence shown in the GSeqEdit report is identical to that of SEQ ID NO: 130 disclosed in the above-identified application.
14. The first 26 amino acid residues of the PRO1244 polypeptide (SEQ ID NO:130) encoded by the cDNA (DNA-64883) are also shown on page 1 of the GSeqEdit report and the remaining 309 residues appear on pages 2-6 of the report.
15. Copies of the pages from the GenenGenes database which report the homology data for the PRO1244 polypeptide sequence are attached to this declaration (with the dates and legal status redacted) as Exhibit B.
16. All activities listed under paragraphs 4-15 were completed prior to June 4, 1998. (See Exhibits A and B).
17. We hereby declare that all statements made herein of our own knowledge are true and that all statements made on information or belief are believed to be true, and further that these statements were made with the knowledge that willful false statements and the like so made are punishable by fine or imprisonment, or both, under Section 1001 of Title 18 of the United States Code and that such willful statements may jeopardize the validity of the application or any patent issued thereon.

  
\_\_\_\_\_  
Napoleone Ferrara, Ph.D.

10/4/04  
\_\_\_\_\_  
Date

\_\_\_\_\_  
Audrey Goddard, Ph.D.

\_\_\_\_\_  
Date

\_\_\_\_\_  
Paul J. Godowski, Ph.D.

\_\_\_\_\_  
Date

\_\_\_\_\_  
James Pan, Ph.D.

\_\_\_\_\_  
Date

\_\_\_\_\_  
Austin Gurney, Ph.D.

\_\_\_\_\_  
Date

\_\_\_\_\_  
Colin K. Watanabe

\_\_\_\_\_  
Date

\_\_\_\_\_  
William I. Wood, Ph.D.

\_\_\_\_\_  
Date

SV 2062079 v1  
10/1/04 2:03 PM (39780.2830)



IN THE UNITED STATES PATENT AND TRADEMARK OFFICE

Applicants: Baker et al. Docket No: 39780-2830P1C8  
Serial No: 10/006,041 Group Art Unit: 1647  
Filed: December 06, 2001 Examiner: Rachel K. Hunnicutt  
For: **SECRETED AND TRANSMEMBRANE POLYPEPTIDES AND NUCLEIC  
ACIDS ENCODING THE SAME**

Commissioner for Patents  
P.O. Box 1450  
Alexandria, Virginia 22313-1450

**DECLARATION OF NAPOLEONE FERRARA, Ph.D.,**  
**AUDREY GODDARD, Ph.D., PAUL J. GODOWSKI, Ph.D.,**  
**AUSTIN GURNEY, Ph.D., JAMES PAN, Ph.D., COLIN K. WATANABE and**  
**WILLIAM I. WOOD, Ph.D. UNDER 37 CFR 1.131**

We, Napoleone Ferrara, Ph.D., Audrey Goddard, Ph.D., Paul J. Godowski, Ph.D., Austin Gurney, Ph.D., James Pan, Ph.D., Colin K. Watanabe and William I. Wood, Ph.D. declare and say as follows:

1. We are the inventors of the above-identified application.
2. We have read and understood the claims pending in this application, and are aware that the claims have been rejected as anticipated by U.S. Patent No. 6,525,174 (Young *et al.*, issue date February 25, 2003 and effective filing date June 4, 1998) and U.S. Patent Publication No. 2003/0096951 (Jacobs *et al.*, publication date May 22, 2003 and effective filing date August 14, 1998).
3. The polypeptide designated as PRO1244 (SEQ ID NO:130) claimed in the above-identified application in the United States was sequenced, cloned and homology to implantation-associated protein identified prior to June 4, 1998.
4. At the time the PRO1244 polypeptide was cloned and sequenced, one of the inventors, Austin Gurney, Ph.D., was responsible for overseeing the cloning of cDNAs which

encoded novel polypeptides, including the cDNA that encoded PRO1244 polypeptide (SEQ ID NO:130) claimed in the above-identified application.

5. At the time the PRO1244 polypeptide was cloned and sequenced, one of the inventors, Audrey Goddard, Ph.D., was, and still is, responsible for overseeing the sequencing of novel polypeptides, including the PRO1244 polypeptide (SEQ ID NO:130) claimed in the above-identified application.
6. At the time the PRO1244 polypeptide was cloned and sequenced, one of the inventors, William I. Wood, was, and still is, responsible for overseeing the homology searches for the novel polypeptides, including that for the PRO1244 polypeptide (SEQ ID NO:130) encoded by the cDNA claimed in the above-identified application.
7. A cDNA clone, referred to as DNA64883-1526 in the above-identified application, was identified as encoding the PRO1244 polypeptide.
8. The full length of the cDNA clone is shown in Figure 73 of the above-identified application. The full-length cDNA sequence has 2213 nucleotide residues. The full length of the PRO1244 peptide encoded by DNA64883-1526 is shown in Figure 74 of the above-identified application. The full-length PRO1244 polypeptide has 335 amino acid residues.
9. Copies of the pages from the GSeqEdit database which report the cloning and sequencing data for the PRO1244 polypeptide sequence and its encoding nucleic acid sequence are attached to this declaration (with the dates redacted) as Exhibit A.
10. The GSeqEdit report shows the full-length nucleic acid sequence for DNA-64883-1526 (identified as "DNA-64883") and the full-length PRO1244 polypeptide encoded by DNA 64883. Both the DNA-64883 and the PRO1244 polypeptide sequences were obtained prior to June 4, 1998.
11. The DNA-64883 sequence shown in the GSeqEdit report is identical to that of SEQ ID NO: 129 disclosed in the above-identified application.
12. The beginning of the cDNA sequence corresponding to SEQ ID NO: 129 in the above-identified application is shown on page 1 of the GSeqEdit database report and

the location of the first nucleotide is marked with "insert starts here" and an arrow. The location of the last nucleotide corresponding to SEQ ID NO: 129 is shown on page 11 and is marked with an arrow.

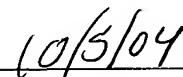
13. The amino acid sequence shown in the GSeqEdit report is identical to that of SEQ ID NO: 130 disclosed in the above-identified application.
14. The first 26 amino acid residues of the PRO1244 polypeptide (SEQ ID NO:130) encoded by the cDNA (DNA-64883) are also shown on page 1 of the GSeqEdit report and the remaining 309 residues appear on pages 2-6 of the report.
15. Copies of the pages from the GenenGenes database which report the homology data for the PRO1244 polypeptide sequence are attached to this declaration (with the dates and legal status redacted) as Exhibit B.
16. All activities listed under paragraphs 4-15 were completed prior to June 4, 1998. (See Exhibits A and B).
17. We hereby declare that all statements made herein of our own knowledge are true and that all statements made on information or belief are believed to be true, and further that these statements were made with the knowledge that willful false statements and the like so made are punishable by fine or imprisonment, or both, under Section 1001 of Title 18 of the United States Code and that such willful statements may jeopardize the validity of the application or any patent issued thereon.

\_\_\_\_\_  
Napoleone Ferrara, Ph.D.



\_\_\_\_\_  
Audrey Goddard, Ph.D.

\_\_\_\_\_  
Date



\_\_\_\_\_  
Date

\_\_\_\_\_  
Paul J. Godowski, Ph.D.

\_\_\_\_\_  
Date

\_\_\_\_\_  
James Pan, Ph.D.

\_\_\_\_\_  
Date

---

Austin Gurney, Ph.D.

---

Date

---

Colin K. Watanabe

---

Date

---

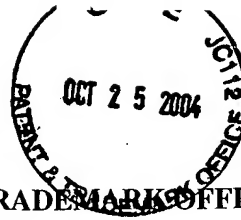
William I. Wood, Ph.D.

---

Date

SV 2062079 v1  
10/1/04 2:03 PM (39780.2830)





IN THE UNITED STATES PATENT AND TRADEMARK OFFICE

Applicants: Baker et al. Docket No: 39780-2830P1C8  
Serial No: 10/006,041 Group Art Unit: 1647  
Filed: December 06, 2001 Examiner: Rachel K. Hunnicutt  
For: **SECRETED AND TRANSMEMBRANE POLYPEPTIDES AND NUCLEIC  
ACIDS ENCODING THE SAME**

Commissioner for Patents  
P.O. Box 1450  
Alexandria, Virginia 22313-1450

**DECLARATION OF NAPOLEONE FERRARA, Ph.D.,**  
**AUDREY GODDARD, Ph.D., PAUL J. GODOWSKI, Ph.D.,**  
**AUSTIN GURNEY, Ph.D., JAMES PAN, Ph.D., COLIN K. WATANABE and**  
**WILLIAM I. WOOD, Ph.D. UNDER 37 CFR 1.131**

We, Napoleone Ferrara, Ph.D., Audrey Goddard, Ph.D., Paul J. Godowski, Ph.D., Austin Gurney, Ph.D., James Pan, Ph.D., Colin K. Watanabe and William I. Wood, Ph.D. declare and say as follows:

1. We are the inventors of the above-identified application.
2. We have read and understood the claims pending in this application, and are aware that the claims have been rejected as anticipated by U.S. Patent No. 6,525,174 (Young *et al.*, issue date February 25, 2003 and effective filing date June 4, 1998) and U.S. Patent Publication No. 2003/0096951 (Jacobs *et al.*, publication date May 22, 2003 and effective filing date August 14, 1998).
3. The polypeptide designated as PRO1244 (SEQ ID NO:130) claimed in the above-identified application in the United States was sequenced, cloned and homology to implantation-associated protein identified prior to June 4, 1998.
4. At the time the PRO1244 polypeptide was cloned and sequenced, one of the inventors, Austin Gurney, Ph.D., was responsible for overseeing the cloning of cDNAs which

encoded novel polypeptides, including the cDNA that encoded PRO1244 polypeptide (SEQ ID NO:130) claimed in the above-identified application.

5. At the time the PRO1244 polypeptide was cloned and sequenced, one of the inventors, Audrey Goddard, Ph.D., was, and still is, responsible for overseeing the sequencing of novel polypeptides, including the PRO1244 polypeptide (SEQ ID NO:130) claimed in the above-identified application.
6. At the time the PRO1244 polypeptide was cloned and sequenced, one of the inventors, William I. Wood, was, and still is, responsible for overseeing the homology searches for the novel polypeptides, including that for the PRO1244 polypeptide (SEQ ID NO:130) encoded by the cDNA claimed in the above-identified application.
7. A cDNA clone, referred to as DNA64883-1526 in the above-identified application, was identified as encoding the PRO1244 polypeptide.
8. The full length of the cDNA clone is shown in Figure 73 of the above-identified application. The full-length cDNA sequence has 2213 nucleotide residues. The full length of the PRO1244 peptide encoded by DNA64883-1526 is shown in Figure 74 of the above-identified application. The full-length PRO1244 polypeptide has 335 amino acid residues.
9. Copies of the pages from the GSeqEdit database which report the cloning and sequencing data for the PRO1244 polypeptide sequence and its encoding nucleic acid sequence are attached to this declaration (with the dates redacted) as Exhibit A.
10. The GSeqEdit report shows the full-length nucleic acid sequence for DNA-64883-1526 (identified as "DNA-64883") and the full-length PRO1244 polypeptide encoded by DNA 64883. Both the DNA-64883 and the PRO1244 polypeptide sequences were obtained prior to June 4, 1998.
11. The DNA-64883 sequence shown in the GSeqEdit report is identical to that of SEQ ID NO: 129 disclosed in the above-identified application.
12. The beginning of the cDNA sequence corresponding to SEQ ID NO: 129 in the above-identified application is shown on page 1 of the GSeqEdit database report and

the location of the first nucleotide is marked with "insert starts here" and an arrow. The location of the last nucleotide corresponding to SEQ ID NO: 129 is shown on page 11 and is marked with an arrow.

13. The amino acid sequence shown in the GSeqEdit report is identical to that of SEQ ID NO: 130 disclosed in the above-identified application.
14. The first 26 amino acid residues of the PRO1244 polypeptide (SEQ ID NO:130) encoded by the cDNA (DNA-64883) are also shown on page 1 of the GSeqEdit report and the remaining 309 residues appear on pages 2-6 of the report.
15. Copies of the pages from the GenenGenes database which report the homology data for the PRO1244 polypeptide sequence are attached to this declaration (with the dates and legal status redacted) as Exhibit B.
16. All activities listed under paragraphs 4-15 were completed prior to June 4, 1998. (See Exhibits A and B).
17. We hereby declare that all statements made herein of our own knowledge are true and that all statements made on information or belief are believed to be true, and further that these statements were made with the knowledge that willful false statements and the like so made are punishable by fine or imprisonment, or both, under Section 1001 of Title 18 of the United States Code and that such willful statements may jeopardize the validity of the application or any patent issued thereon.

\_\_\_\_\_  
Napoleone Ferrara, Ph.D.

\_\_\_\_\_  
Date

\_\_\_\_\_  
Andrey Goddard, Ph.D.

\_\_\_\_\_  
Date

  
\_\_\_\_\_  
Paul J. Godowski, Ph.D.

10/05/01  
\_\_\_\_\_  
Date

\_\_\_\_\_  
James Pan, Ph.D.

\_\_\_\_\_  
Date

---

Austin Gurney, Ph.D.

---

Date

---

Colin K. Watanabe

---

Date

---

William I. Wood, Ph.D.

---

Date

SV 2062079 v1  
10/1/04 2:03 PM (39780.2830)



IN THE UNITED STATES PATENT AND TRADEMARK OFFICE

Applicants: Baker et al. Docket No: 39780-2830P1C8  
Serial No: 10/006,041 Group Art Unit: 1647  
Filed: December 06, 2001 Examiner: Rachel K. Hunnicutt  
For: **SECRETED AND TRANSMEMBRANE POLYPEPTIDES AND NUCLEIC  
ACIDS ENCODING THE SAME**

Commissioner for Patents  
P.O. Box 1450  
Alexandria, Virginia 22313-1450

**DECLARATION OF NAPOLEONE FERRARA, Ph.D.,**  
**AUDREY GODDARD, Ph.D., PAUL J. GODOWSKI, Ph.D.,**  
**AUSTIN GURNEY, Ph.D., JAMES PAN, Ph.D., COLIN K. WATANABE and**  
**WILLIAM I. WOOD, Ph.D. UNDER 37 CFR 1.131**

We, Napoleone Ferrara, Ph.D., Audrey Goddard, Ph.D., Paul J. Godowski, Ph.D., Austin Gurney, Ph.D., James Pan, Ph.D., Colin K. Watanabe and William I. Wood, Ph.D. declare and say as follows:

1. We are the inventors of the above-identified application.
2. We have read and understood the claims pending in this application, and are aware that the claims have been rejected as anticipated by U.S. Patent No. 6,525,174 (Young *et al.*, issue date February 25, 2003 and effective filing date June 4, 1998) and U.S. Patent Publication No. 2003/0096951 (Jacobs *et al.*, publication date May 22, 2003 and effective filing date August 14, 1998).
3. The polypeptide designated as PRO1244 (SEQ ID NO:130) claimed in the above-identified application in the United States was sequenced, cloned and homology to implantation-associated protein identified prior to June 4, 1998.
4. At the time the PRO1244 polypeptide was cloned and sequenced, one of the inventors, Austin Gurney, Ph.D., was responsible for overseeing the cloning of cDNAs which

encoded novel polypeptides, including the cDNA that encoded PRO1244 polypeptide (SEQ ID NO:130) claimed in the above-identified application.

5. At the time the PRO1244 polypeptide was cloned and sequenced, one of the inventors, Audrey Goddard, Ph.D., was, and still is, responsible for overseeing the sequencing of novel polypeptides, including the PRO1244 polypeptide (SEQ ID NO:130) claimed in the above-identified application.
6. At the time the PRO1244 polypeptide was cloned and sequenced, one of the inventors, William I. Wood, was, and still is, responsible for overseeing the homology searches for the novel polypeptides, including that for the PRO1244 polypeptide (SEQ ID NO:130) encoded by the cDNA claimed in the above-identified application.
7. A cDNA clone, referred to as DNA64883-1526 in the above-identified application, was identified as encoding the PRO1244 polypeptide.
8. The full length of the cDNA clone is shown in Figure 73 of the above-identified application. The full-length cDNA sequence has 2213 nucleotide residues. The full length of the PRO1244 peptide encoded by DNA64883-1526 is shown in Figure 74 of the above-identified application. The full-length PRO1244 polypeptide has 335 amino acid residues.
9. Copies of the pages from the GSeqEdit database which report the cloning and sequencing data for the PRO1244 polypeptide sequence and its encoding nucleic acid sequence are attached to this declaration (with the dates redacted) as Exhibit A.
10. The GSeqEdit report shows the full-length nucleic acid sequence for DNA-64883-1526 (identified as "DNA-64883") and the full-length PRO1244 polypeptide encoded by DNA 64883. Both the DNA-64883 and the PRO1244 polypeptide sequences were obtained prior to June 4, 1998.
11. The DNA-64883 sequence shown in the GSeqEdit report is identical to that of SEQ ID NO: 129 disclosed in the above-identified application.
12. The beginning of the cDNA sequence corresponding to SEQ ID NO: 129 in the above-identified application is shown on page 1 of the GSeqEdit database report and

the location of the first nucleotide is marked with "insert starts here" and an arrow. The location of the last nucleotide corresponding to SEQ ID NO: 129 is shown on page 11 and is marked with an arrow.

13. The amino acid sequence shown in the GSeqEdit report is identical to that of SEQ ID NO: 130 disclosed in the above-identified application.
14. The first 26 amino acid residues of the PRO1244 polypeptide (SEQ ID NO: 130) encoded by the cDNA (DNA-64883) are also shown on page 1 of the GSeqEdit report and the remaining 309 residues appear on pages 2-6 of the report.
15. Copies of the pages from the GenenGenes database which report the homology data for the PRO1244 polypeptide sequence are attached to this declaration (with the dates and legal status redacted) as Exhibit B.
16. All activities listed under paragraphs 4-15 were completed prior to June 4, 1998. (See Exhibits A and B).
17. We hereby declare that all statements made herein of our own knowledge are true and that all statements made on information or belief are believed to be true, and further that these statements were made with the knowledge that willful false statements and the like so made are punishable by fine or imprisonment, or both, under Section 1001 of Title 18 of the United States Code and that such willful statements may jeopardize the validity of the application or any patent issued thereon.

\_\_\_\_\_  
Napoleone Ferrara, Ph.D.

\_\_\_\_\_  
Date

\_\_\_\_\_  
Audrey Goddard, Ph.D.

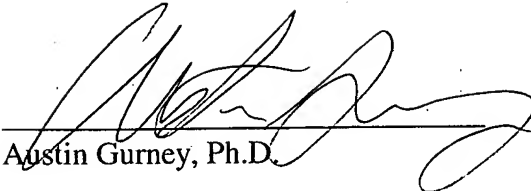
\_\_\_\_\_  
Date

\_\_\_\_\_  
Paul J. Godowski, Ph.D.

\_\_\_\_\_  
Date

\_\_\_\_\_  
James Pan, Ph.D.

\_\_\_\_\_  
Date

  
Austin Gurney, Ph.D.

10/5/04  
Date

\_\_\_\_\_  
Colin K. Watanabe

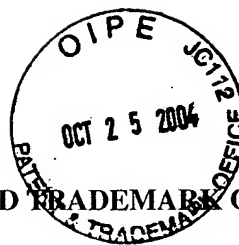
\_\_\_\_\_  
Date

\_\_\_\_\_  
William I. Wood, Ph.D.

\_\_\_\_\_  
Date

SV 2062079 v1  
10/1/04 2:03 PM (39780.2830)





IN THE UNITED STATES PATENT AND TRADEMARK OFFICE

Applicants: Baker et al. Docket No: 39780-2830P1C8  
Serial No: 10/006,041 Group Art Unit: 1647  
Filed: December 06, 2001 Examiner: Rachel K. Hunnicutt  
For: **SECRETED AND TRANSMEMBRANE POLYPEPTIDES AND NUCLEIC  
ACIDS ENCODING THE SAME**

Commissioner for Patents  
P.O. Box 1450  
Alexandria, Virginia 22313-1450

**DECLARATION OF NAPOLEONE FERRARA, Ph.D.,**  
**AUDREY GODDARD, Ph.D., PAUL J. GODOWSKI, Ph.D.,**  
**AUSTIN GURNEY, Ph.D., JAMES PAN, Ph.D., COLIN K. WATANABE and**  
**WILLIAM I. WOOD, Ph.D. UNDER 37 CFR 1.131**

We, Napoleone Ferrara, Ph.D., Audrey Goddard, Ph.D., Paul J. Godowski, Ph.D., Austin Gurney, Ph.D., James Pan, Ph.D., Colin K. Watanabe and William I. Wood, Ph.D. declare and say as follows:

1. We are the inventors of the above-identified application.
2. We have read and understood the claims pending in this application, and are aware that the claims have been rejected as anticipated by U.S. Patent No. 6,525,174 (Young *et al.*, issue date February 25, 2003 and effective filing date June 4, 1998) and U.S. Patent Publication No. 2003/0096951 (Jacobs *et al.*, publication date May 22, 2003 and effective filing date August 14, 1998).
3. The polypeptide designated as PRO1244 (SEQ ID NO:130) claimed in the above-identified application in the United States was sequenced, cloned and homology to implantation-associated protein identified prior to June 4, 1998.
4. At the time the PRO1244 polypeptide was cloned and sequenced, one of the inventors, Austin Gurney, Ph.D., was responsible for overseeing the cloning of cDNAs which

encoded novel polypeptides, including the cDNA that encoded PRO1244 polypeptide (SEQ ID NO:130) claimed in the above-identified application.

5. At the time the PRO1244 polypeptide was cloned and sequenced, one of the inventors, Audrey Goddard, Ph.D., was, and still is, responsible for overseeing the sequencing of novel polypeptides, including the PRO1244 polypeptide (SEQ ID NO:130) claimed in the above-identified application.
6. At the time the PRO1244 polypeptide was cloned and sequenced, one of the inventors, William I. Wood, was, and still is, responsible for overseeing the homology searches for the novel polypeptides, including that for the PRO1244 polypeptide (SEQ ID NO:130) encoded by the cDNA claimed in the above-identified application.
7. A cDNA clone, referred to as DNA64883-1526 in the above-identified application, was identified as encoding the PRO1244 polypeptide.
8. The full length of the cDNA clone is shown in Figure 73 of the above-identified application. The full-length cDNA sequence has 2213 nucleotide residues. The full length of the PRO1244 peptide encoded by DNA64883-1526 is shown in Figure 74 of the above-identified application. The full-length PRO1244 polypeptide has 335 amino acid residues.
9. Copies of the pages from the GSeqEdit database which report the cloning and sequencing data for the PRO1244 polypeptide sequence and its encoding nucleic acid sequence are attached to this declaration (with the dates redacted) as Exhibit A.
10. The GSeqEdit report shows the full-length nucleic acid sequence for DNA-64883-1526 (identified as "DNA-64883") and the full-length PRO1244 polypeptide encoded by DNA 64883. Both the DNA-64883 and the PRO1244 polypeptide sequences were obtained prior to June 4, 1998.
11. The DNA-64883 sequence shown in the GSeqEdit report is identical to that of SEQ ID NO: 129 disclosed in the above-identified application.
12. The beginning of the cDNA sequence corresponding to SEQ ID NO: 129 in the above-identified application is shown on page 1 of the GSeqEdit database report and

the location of the first nucleotide is marked with "insert starts here" and an arrow. The location of the last nucleotide corresponding to SEQ ID NO: 129 is shown on page 11 and is marked with an arrow.

13. The amino acid sequence shown in the GSeqEdit report is identical to that of SEQ ID NO: 130 disclosed in the above-identified application.
14. The first 26 amino acid residues of the PRO1244 polypeptide (SEQ ID NO:130) encoded by the cDNA (DNA-64883) are also shown on page 1 of the GSeqEdit report and the remaining 309 residues appear on pages 2-6 of the report.
15. Copies of the pages from the GenenGenes database which report the homology data for the PRO1244 polypeptide sequence are attached to this declaration (with the dates and legal status redacted) as Exhibit B.
16. All activities listed under paragraphs 4-15 were completed prior to June 4, 1998. (See Exhibits A and B).
17. We hereby declare that all statements made herein of our own knowledge are true and that all statements made on information or belief are believed to be true, and further that these statements were made with the knowledge that willful false statements and the like so made are punishable by fine or imprisonment, or both, under Section 1001 of Title 18 of the United States Code and that such willful statements may jeopardize the validity of the application or any patent issued thereon.

\_\_\_\_\_  
Napoleone Ferrara, Ph.D.

\_\_\_\_\_  
Date

\_\_\_\_\_  
Audrey Goddard, Ph.D.

\_\_\_\_\_  
Date

\_\_\_\_\_  
Paul J. Godowski, Ph.D.

\_\_\_\_\_  
Date

\_\_\_\_\_  
James Pan, Ph.D.

\_\_\_\_\_  
Date

*Oct 08 / 2004*

---

Austin Gurney, Ph.D.

---

Date

---

Colin K. Watanabe

---

Date

---

William I. Wood, Ph.D.

---

Date

SV 2062079 v1  
10/1/04 2:03 PM (39780.2830)



IN THE UNITED STATES PATENT AND TRADEMARK OFFICE

Applicants: Baker et al. Docket No: 39780-2830P1C8  
Serial No: 10/006,041 Group Art Unit: 1647  
Filed: December 06, 2001 Examiner: Rachel K. Hunnicutt  
For: **SECRETED AND TRANSMEMBRANE POLYPEPTIDES AND NUCLEIC  
ACIDS ENCODING THE SAME**

Commissioner for Patents  
P.O. Box 1450  
Alexandria, Virginia 22313-1450

**DECLARATION OF NAPOLEONE FERRARA, Ph.D.,**  
**AUDREY GODDARD, Ph.D., PAUL J. GODOWSKI, Ph.D.,**  
**AUSTIN GURNEY, Ph.D., JAMES PAN, Ph.D., COLIN K. WATANABE and**  
**WILLIAM I. WOOD, Ph.D. UNDER 37 CFR 1.131**

We, Napoleone Ferrara, Ph.D., Audrey Goddard, Ph.D., Paul J. Godowski, Ph.D., Austin Gurney, Ph.D., James Pan, Ph.D., Colin K. Watanabe and William I. Wood, Ph.D. declare and say as follows:

1. We are the inventors of the above-identified application.
2. We have read and understood the claims pending in this application, and are aware that the claims have been rejected as anticipated by U.S. Patent No. 6,525,174 (Young *et al.*, issue date February 25, 2003 and effective filing date June 4, 1998) and U.S. Patent Publication No. 2003/0096951 (Jacobs *et al.*, publication date May 22, 2003 and effective filing date August 14, 1998).
3. The polypeptide designated as PRO1244 (SEQ ID NO:130) claimed in the above-identified application in the United States was sequenced, cloned and homology to implantation-associated protein identified prior to June 4, 1998.
4. At the time the PRO1244 polypeptide was cloned and sequenced, one of the inventors, Austin Gurney, Ph.D., was responsible for overseeing the cloning of cDNAs which

encoded novel polypeptides, including the cDNA that encoded PRO1244 polypeptide (SEQ ID NO:130) claimed in the above-identified application.

5. At the time the PRO1244 polypeptide was cloned and sequenced, one of the inventors, Audrey Goddard, Ph.D., was, and still is, responsible for overseeing the sequencing of novel polypeptides, including the PRO1244 polypeptide (SEQ ID NO:130) claimed in the above-identified application.
6. At the time the PRO1244 polypeptide was cloned and sequenced, one of the inventors, William I. Wood, was, and still is, responsible for overseeing the homology searches for the novel polypeptides, including that for the PRO1244 polypeptide (SEQ ID NO:130) encoded by the cDNA claimed in the above-identified application.
7. A cDNA clone, referred to as DNA64883-1526 in the above-identified application, was identified as encoding the PRO1244 polypeptide.
8. The full length of the cDNA clone is shown in Figure 73 of the above-identified application. The full-length cDNA sequence has 2213 nucleotide residues. The full length of the PRO1244 peptide encoded by DNA64883-1526 is shown in Figure 74 of the above-identified application. The full-length PRO1244 polypeptide has 335 amino acid residues.
9. Copies of the pages from the GSeqEdit database which report the cloning and sequencing data for the PRO1244 polypeptide sequence and its encoding nucleic acid sequence are attached to this declaration (with the dates redacted) as Exhibit A.
10. The GSeqEdit report shows the full-length nucleic acid sequence for DNA-64883-1526 (identified as "DNA-64883") and the full-length PRO1244 polypeptide encoded by DNA 64883. Both the DNA-64883 and the PRO1244 polypeptide sequences were obtained prior to June 4, 1998.
11. The DNA-64883 sequence shown in the GSeqEdit report is identical to that of SEQ ID NO: 129 disclosed in the above-identified application.
12. The beginning of the cDNA sequence corresponding to SEQ ID NO: 129 in the above-identified application is shown on page 1 of the GSeqEdit database report and

the location of the first nucleotide is marked with "insert starts here" and an arrow. The location of the last nucleotide corresponding to SEQ ID NO: 129 is shown on page 11 and is marked with an arrow.

13. The amino acid sequence shown in the GSeqEdit report is identical to that of SEQ ID NO: 130 disclosed in the above-identified application.
14. The first 26 amino acid residues of the PRO1244 polypeptide (SEQ ID NO:130) encoded by the cDNA (DNA-64883) are also shown on page 1 of the GSeqEdit report and the remaining 309 residues appear on pages 2-6 of the report.
15. Copies of the pages from the GenenGenes database which report the homology data for the PRO1244 polypeptide sequence are attached to this declaration (with the dates and legal status redacted) as Exhibit B.
16. All activities listed under paragraphs 4-15 were completed prior to June 4, 1998. (See Exhibits A and B).
17. We hereby declare that all statements made herein of our own knowledge are true and that all statements made on information or belief are believed to be true, and further that these statements were made with the knowledge that willful false statements and the like so made are punishable by fine or imprisonment, or both, under Section 1001 of Title 18 of the United States Code and that such willful statements may jeopardize the validity of the application or any patent issued thereon.

\_\_\_\_\_  
Napoleone Ferrara, Ph.D.

\_\_\_\_\_  
Date

\_\_\_\_\_  
Audrey Goddard, Ph.D.

\_\_\_\_\_  
Date

\_\_\_\_\_  
Paul J. Godowski, Ph.D.

\_\_\_\_\_  
Date

\_\_\_\_\_  
James Pan, Ph.D.

\_\_\_\_\_  
Date

\_\_\_\_\_  
Austin Gurney, Ph.D.

Colin K. Watanabe  
Colin K. Watanabe

\_\_\_\_\_  
Date

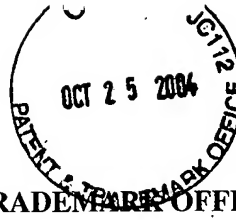
Oct 8, 2004  
Date

\_\_\_\_\_  
William I. Wood, Ph.D.

\_\_\_\_\_  
Date

SV 2062079 v1  
10/1/04 2:03 PM (39780.2830)





IN THE UNITED STATES PATENT AND TRADEMARK OFFICE

Applicants: Baker et al. Docket No: 39780-2830P1C8  
Serial No: 10/006,041 Group Art Unit: 1647  
Filed: December 06, 2001 Examiner: Rachel K. Hunnicutt  
For: **SECRETED AND TRANSMEMBRANE POLYPEPTIDES AND NUCLEIC  
ACIDS ENCODING THE SAME**

Commissioner for Patents  
P.O. Box 1450  
Alexandria, Virginia 22313-1450

**DECLARATION OF NAPOLEONE FERRARA, Ph.D.,**  
**AUDREY GODDARD, Ph.D., PAUL J. GODOWSKI, Ph.D.,**  
**AUSTIN GURNEY, Ph.D., JAMES PAN, Ph.D., COLIN K. WATANABE and**  
**WILLIAM I. WOOD, Ph.D. UNDER 37 CFR 1.131**

We, Napoleone Ferrara, Ph.D., Audrey Goddard, Ph.D., Paul J. Godowski, Ph.D., Austin Gurney, Ph.D., James Pan, Ph.D., Colin K. Watanabe and William I. Wood, Ph.D. declare and say as follows:

1. We are the inventors of the above-identified application.
2. We have read and understood the claims pending in this application, and are aware that the claims have been rejected as anticipated by U.S. Patent No. 6,525,174 (Young *et al.*, issue date February 25, 2003 and effective filing date June 4, 1998) and U.S. Patent Publication No. 2003/0096951 (Jacobs *et al.*, publication date May 22, 2003 and effective filing date August 14, 1998).
3. The polypeptide designated as PRO1244 (SEQ ID NO:130) claimed in the above-identified application in the United States was sequenced, cloned and homology to implantation-associated protein identified prior to June 4, 1998.
4. At the time the PRO1244 polypeptide was cloned and sequenced, one of the inventors, Austin Gurney, Ph.D., was responsible for overseeing the cloning of cDNAs which

encoded novel polypeptides, including the cDNA that encoded PRO1244 polypeptide (SEQ ID NO:130) claimed in the above-identified application.

5. At the time the PRO1244 polypeptide was cloned and sequenced, one of the inventors, Audrey Goddard, Ph.D., was, and still is, responsible for overseeing the sequencing of novel polypeptides, including the PRO1244 polypeptide (SEQ ID NO:130) claimed in the above-identified application.
6. At the time the PRO1244 polypeptide was cloned and sequenced, one of the inventors, William I. Wood, was, and still is, responsible for overseeing the homology searches for the novel polypeptides, including that for the PRO1244 polypeptide (SEQ ID NO:130) encoded by the cDNA claimed in the above-identified application.
7. A cDNA clone, referred to as DNA64883-1526 in the above-identified application, was identified as encoding the PRO1244 polypeptide.
8. The full length of the cDNA clone is shown in Figure 73 of the above-identified application. The full-length cDNA sequence has 2213 nucleotide residues. The full length of the PRO1244 peptide encoded by DNA64883-1526 is shown in Figure 74 of the above-identified application. The full-length PRO1244 polypeptide has 335 amino acid residues.
9. Copies of the pages from the GSeqEdit database which report the cloning and sequencing data for the PRO1244 polypeptide sequence and its encoding nucleic acid sequence are attached to this declaration (with the dates redacted) as Exhibit A.
10. The GSeqEdit report shows the full-length nucleic acid sequence for DNA-64883-1526 (identified as "DNA-64883") and the full-length PRO1244 polypeptide encoded by DNA 64883. Both the DNA-64883 and the PRO1244 polypeptide sequences were obtained prior to June 4, 1998.
11. The DNA-64883 sequence shown in the GSeqEdit report is identical to that of SEQ ID NO: 129 disclosed in the above-identified application.
12. The beginning of the cDNA sequence corresponding to SEQ ID NO: 129 in the above-identified application is shown on page 1 of the GSeqEdit database report and

the location of the first nucleotide is marked with "insert starts here" and an arrow. The location of the last nucleotide corresponding to SEQ ID NO: 129 is shown on page 11 and is marked with an arrow.

13. The amino acid sequence shown in the GSeqEdit report is identical to that of SEQ ID NO: 130 disclosed in the above-identified application.
14. The first 26 amino acid residues of the PRO1244 polypeptide (SEQ ID NO:130) encoded by the cDNA (DNA-64883) are also shown on page 1 of the GSeqEdit report and the remaining 309 residues appear on pages 2-6 of the report.
15. Copies of the pages from the GenenGenes database which report the homology data for the PRO1244 polypeptide sequence are attached to this declaration (with the dates and legal status redacted) as Exhibit B.
16. All activities listed under paragraphs 4-15 were completed prior to June 4, 1998. (See Exhibits A and B).
17. We hereby declare that all statements made herein of our own knowledge are true and that all statements made on information or belief are believed to be true, and further that these statements were made with the knowledge that willful false statements and the like so made are punishable by fine or imprisonment, or both, under Section 1001 of Title 18 of the United States Code and that such willful statements may jeopardize the validity of the application or any patent issued thereon.

\_\_\_\_\_  
Napoleone Ferrara, Ph.D.

\_\_\_\_\_  
Date

\_\_\_\_\_  
Audrey Goddard, Ph.D.

\_\_\_\_\_  
Date

\_\_\_\_\_  
Paul J. Godowski, Ph.D.

\_\_\_\_\_  
Date

\_\_\_\_\_  
James Pan, Ph.D.

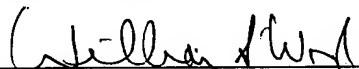
\_\_\_\_\_  
Date

\_\_\_\_\_  
Austin Gurney, Ph.D.

\_\_\_\_\_  
Date

\_\_\_\_\_  
Colin K. Watanabe

\_\_\_\_\_  
Date

\_\_\_\_\_  
  
William I. Wood, Ph.D.

\_\_\_\_\_  
10/5/04  
Date

SV 2062079 v1  
10/1/04 2:03 PM (39780.2830)

> [REDACTED]  
 >DNA64883 [Full]  
 >510 Sites [All Sites]  
 > [REDACTED] DNA64883 wlv GSeqEdit  
 > [REDACTED] DNA64883 zemin GSeqEdit  
 > [REDACTED] DNA64883 goddarda GSeqEdit  
 > [REDACTED] DNA64883 sheldens GSeqEdit  
 >HBN64883.seq, sequenced at ABI/ACGT by Peter Ma and Eilson Chen  
 >human ortholog of implantation-associated protein - Rattus

mulI		nlaIII
tagI		msII
xhoI	thai	styI
cllI	fnuDI/mvnI	ncol
smlI	hinfI	dsai
paer7I	acII	tseI
tsp509I[M.ecoRI-]	tseI bstuI[M.hai-]	btgl/bstDSI
ecoRI	fnu4HI/bsofI	fnu4HI/bsofI
apoI	bvi bsh1236I	bstXI
nlaIII	bvi bsh1236I	bvII
hhaI/cfoI	tsp45I	hinfI
bsmAI	maeIII	hhaI/cfoI
haeII	hpy99I	maeII/hpyC
hpy99I	mulI	

1 CGGAATTTCGG CTCGAGGAGC GAACATGGCA GCGCGTTGGC GGTTCGTGTG TGTCTCTGTG ACCATGCTGG TGGCGCTGCT CATCGTTTGC GACGTTCCCT  
 GCCTTAAGCC GAGCTCCTCG CTGTACCGT CGCGCAACCG CCAAAACAC ACAGAGACAC TGTACCCACC ACCGCGACGA GTAGCAAACG CTGCAAGGGA  
 1 M A A R W R F W C V S V T M V V A L L I V C D V P S  
 ^insert starts here  
 ^MET

	mulI		alul
	alwNI(dcm-)		pvuII
	alw26I/bsmAI	bsaXI	mspAII/nspBII
		hpy188I	
101	CAGCCTCTGC CCAAGAAG AAGGATG TGTATCTGA AAAGTTAGT CAGTGTATG AATGACTAA CAAAGACT GTATTAAGA TGAATGAGA		bsmAI
	GTGGAGACG GGTTCCTTC TTCTCTACC ACAATAGACT TTCCAATCA GTGACTACC TTACCTGATT GTTCTTGA CATTAATCTT ACTTACCTCT		
27	A S A Q R K K E M V L S E K V S Q L M E W T N K R P V I R M N G D		
		tsprI	bst4CI/hpyCH4III
		btsI	ahdI/eam1105I
	hpy99I	nlaiII	hpyCH4V al
		tsp509I	
201	CAAGTTCGT CGCCTTGTGA AAGCCACC GAGAATTAC TCCGTATCG TCATGTTAC TCCTCTCAA CTGCATAGAC AGTGTGCGT TTGCAAGCAA		
	GTCAAGGCA GCGAACACT TTCGGGGTGG CTCTTAAAG AGCAATAGC AGTACAAGTG ACGAGAGTT GACGTATCTG TCACACAGCA AACGTTGCTT		
60	K F R R L V K A P P R N Y S V I V M F T A L Q L H R Q C V V C K Q		

scrFI[dcn-]  
 pspGI  
 mvai  
 ecorII[dcn-]  
 dsav[dcn-]  
 bstNI  
 bssKI[dcn-]  
 apyI[dcn+]  
 sau3AI  
 mboI/ndeII[dam-]  
 dpnII[dam-]  
 dpnI[dam+]  
 alwI[dam-]  
 bstYI/xhoII  
 alwNI[dcn-]  
 alw26I/bsmAI  
 tsp509I[M.ecorI-]  
 ecorI pfIMI[dcn-]  
 apoI bslI[dcn-]  
 mboII hpy188III  
 301 GCTGATGAAG AATTCAGAT CCTGCAAC TCTGGCGAT ACTCCAGTGC ATTCAACCAAC AGGATATTIT TTGCCATGCT GGATTITGAT GAAGGCTCTG  
 CGACTACTTC TTAAAGTCTA GGACCGTTTG AGGACCGCTA TGAGTCAAG TAAGTGTG TTCTATAAAA AAGGTACCA CCTAAACTA CTCCGAGAC  
 93 A D E E F Q I L A N S W R Y S S A F T N R I F F A M V D F D E G S D  
 apyI[dcn+]  
 bssKI[dcn-]  
 bstNI baeI  
 tspRI  
 hpyCH4V  
 nlaIII  
 styI  
 ncoI  
 dsal  
 btgl/bstDSI  
 bsauI  
 hpy18

tsp5091[M.ecori-]

ecori

hpyCH4V

sfani

apoi

econI

hpy188I

nlaiII aluI

bsII

bsII

hphI

ndel

maeiII

acII

401 ATGTAATTCA GATGCTAAC ATGAATTCAG CTCCACTTT CATCACTTT CCTGCAAAAG GGAACCCAA ACCGGGTGAT ACATATGAGT TACAGTGGC  
TACATAAAGT CTACGATTTG TACTTAAGTC GAGGTGAAA GTAGTGAAG GAGCTTTTC CCTTGGGTT TGCCCCACTA TGTATACTCA ATGTCCAGC  
127 Y F Q M L N M N S A P T F I N F P A K G K P K R G D T Y E L Q V R

ddeI[M.aluI-]

bspcNI

mspi

saU3AI

celII/espi

hpaII

mboI/ndeII[dam-]

b1pI/bpu1102I scrFI[M.hpaII-]

aluI

ncII

dpnII[dam-]

pvuII

dsav

dpnI[dam+]

mspAII/nsbII

bskI

alwI[dam-]

spsI

tsp509I

avaII

bsII

501 GGGTTTTC GGTGACGAGA TTGCCCCGTG GATGCCCGAC AGACTGATG TCAATATTAG AGTATTAGA CCCCCAAATT ATGCTGCTCC CCTATGTTG  
CCCAAAAGT CGACTCGCTT AACGGGCCAC CTAGCGGCTG TCTGACTAC AGTATAATC TCACTAATCT GGGGTTTAA TAGACCCAGG GGAATACAAC  
160 G F S A E Q I A R W I A D R T D V N I R V I R P P N Y A G P L M L

taqI

sfuI

bsfBI

bslCI

baeI

mboII mboII

apoi

mseI

bsrI

mwoI hpyCH4V

601 GGATTGCTTT TGGCTGTAT TGGTGAAGCTT GGTATCTTC GAAGAAGTAA TATGAATTT CTCTTATAA AAAGTGAAG GGTCTTGA GCTTGTGTT

CCTAACGAAA ACCGACATA ACCACTGAA CACATAGAG CTCTTCAAT ATACCTTAA GAGAAATTA TTTGACCTAC CCGAAGAGT CGAAGACAAA  
193 G L L L A V I G G L V Y L R R S N M E F L F N K T G W A F A A L C F



nlaIII                      nlaIII  
 pciI                      styI  
 nsphi                      ncoI  
 nspl                      dsal  
 bclI                      bglI/bs  
 tflI                      aflIII  
 hinfI                      bsmFI                      tsp509I                      bsaJI  
 ndel                      cac8I                      ahdI/eam1105I                      nlaIV                      mnlI                      bsa96I  
 avall                     

701 TTGTGCTTGC TATGACATCT GGTCAAATGT GGAACCATAT AAGAGACCA CCATAAGCCC ATAAGAATCC CCACACGGGA CATGTGAATT ATATCCAATGG  
 AACACGAACG ATACTGTAGA CCAGTTTACA CCTTGCTGTA TTCTCCTGCT GGTATACGGG TATTTCTTAGG GGTGTGCCCT GTACACTTAA TATAGTACC  
 227 V L A M T S G Q M W N H I R G P P Y A H K N P H T G H V N Y I H G

ddeI                      tseI  
 eco8II                      fnu4HI/bsoFI  
 tru9I                      bvuI                      hpyI  
 bsu36I/mscII/sauI                      aluI                      mnlI  
 maeIII  
 mboII                      mscI  
 aluI                     

801 AAGCAGTCAA GCCCAGTTTG TAGCTGAAC ACACATTTGT CTCTGTTTA ATGGTGAGT TACCTTAGGA ATGGTGCTTT TATGTGAAC TGCTACCTCT  
 TTCGTCAGTT CGGCTCAAC ATGCACTTTG TGTGTACAA GAAGACAAT TACCACCTCA ATGGAATCCT TACCACGAAA ATACACTTCG ACGATGAGA  
 260 S S Q A Q F V A E T H I V L L F N G G V T L G M V L L C E A A T S

sau3AI  
 mboI/ndelI[dam  
 dpmII[dam-]  
 sfanI                      dpmI[dam+]  
 foki                      bstYI/xhoII  
 bstF5I                      bglII                      nl  
 mboII                     

901 GACATGAGATA TTGGAAGCG AAAGATAATG TGTGTGCTG GATTTGACT TGTGTATTA TTCTTCAGTT GGAATGCTCTC TATTTTAGA TCTAATATAC  
 CTGTACCTAT AACCTTCCG TTCTATTTAC ACACACCGAC CATTAACCTGA ACAACATAT AAGAGTCAA CCTACGAGAG ATAAAATCT AGATTTATAG  
 293 D M D I G K R K I M C V A G I G L V V L F F S W M L S I F R S K Y H

**bsri**

rsai

bpmI/gsuI[dcn-]

**bsrI csp6I**

tsPRI scai tsps091

327. G Y P Y S F L M S O

msei trugI

**ahaii/drai**

# man 1.1

**msel**

msei

II OGPU

msel

TTCTTCTTAC GTTGACATA TTAACATTA AGTTCACCTA AATTATCA TTAGTAATTT GGTTCCTCTT ACACATCACC GAATTGTTGG

**αδεΙ**

1621 TUGI

I tsprti

160509I

trug

hpy188I

tsp509I

earl/ksp6321

msci

csp6I msei

TTAGGAGACA GTTTAGACT CCATAACTT TTATTATAG GAGATTGA AGAGAGGGT CACTGAAAT ACCTTGTA A TTAATCATG TTATTCATA

mset

# h1ncII/h1ndII hpy188I

**alut**

b31I

TAATATTTTT AACATTTTGA TGATGAACA AAATCATCT TGTTCGAGT TTTGATGAA TCAATTGAAC CAGTAGACTA AAATATAACG GAATAGCTTTT

scrFI[dcM-]

pspGI

mvaI

ecorII[dcM-]

dsaV[dcM-]

bstNI

bsaKI[dcM-]

apyI[dcM+]

sexAI

tsp509I[M.ecorI-]

xmnI

ecorI

asp700

apoI

ddel[M.aluI-]

bsf5I

mboII

aluI

msII

foKI

taiI

hgiAI/aspHI

bspI286

bsiHKAI

hpy188I

eco57I

mbolI

bmyI

bfai

muli

hpy188I

mbolI

bpuAI

hpaII

bsi

hpaII

hpy188I

mbolI

bpuAI

hpaII

bsi

hpaII

hpy188I

mbolI

bpuAI

hpaII

bsi

hpaII

hpy188I

mbolI

bpuAI

hpaII

bsi

hpaII

hpy188I

mbolI

bpuAI

hpaII

bsi

hpaII

hpy188I

mbolI

bpuAI

hpaII

bsi

hpaII

hpy188I

mbolI

bpuAI

hpaII

bsi

hpaII

hpy188I

mbolI

bpuAI

hpaII

bsi

hpaII

hpy188I

mbolI

bpuAI

hpaII

bsi

hpaII

hpy188I

mbolI

bpuAI

hpaII

bsi

hpaII

hpy188I

mbolI

bpuAI

hpaII

bsi

hpaII

hpy188I

mbolI

bpuAI

hpaII

bsi

hpaII

hpy188I

mbolI

bpuAI

hpaII

bsi

hpaII

hpy188I

mbolI

bpuAI

hpaII

bsi

hpaII

hpy188I

mbolI

bpuAI

hpaII

bsi

hpaII

hpy188I

mbolI

bpuAI

hpaII

bsi

hpaII

hpy188I

mbolI

bpuAI

hpaII

bsi

hpaII

hpy188I

mbolI

bpuAI

hpaII

bsi

hpaII

hpy188I

mbolI

bpuAI

hpaII

bsi

hpaII

hpy188I

mbolI

bpuAI

hpaII

bsi

hpaII

hpy188I

mbolI

bpuAI

hpaII

bsi

hpaII

hpy188I

mbolI

bpuAI

hpaII

bsi

hpaII

hpy188I

mbolI

bpuAI

hpaII

bsi

hpaII

hpy188I

mbolI

bpuAI

hpaII

bsi

hpaII

hpy188I

mbolI

bpuAI

hpaII

bsi

hpaII

hpy188I

mbolI

bpuAI

hpaII

bsi

hpaII

hpy188I

mbolI

bpuAI

hpaII

bsi

hpaII

hpy188I

mbolI

bpuAI

hpaII

bsi

hpaII

hpy188I

mbolI

bpuAI

hpaII

bsi

hpaII

hpy188I

mbolI

bpuAI

hpaII

bsi

hpaII

hpy188I

mbolI

bpuAI

hpaII

bsi

hpaII

hpy188I

mbolI

bpuAI

hpaII

bsi

hpaII

hpy188I

mbolI

bpuAI

hpaII

bsi

hpaII

hpy188I

mbolI

bpuAI

hpaII

bsi

hpaII

hpy188I

mbolI

bpuAI

hpaII

bsi

hpaII

hpy188I

mbolI

bpuAI

hpaII

bsi

hpaII

hpy188I

mbolI

bpuAI

hpaII

bsi

hpaII

hpy188I

mbolI

bpuAI

hpaII

bsi

hpaII

hpy188I

mbolI

bpuAI

hpaII

bsi

hpaII

hpy188I

mbolI

bpuAI

hpaII

bsi

hpaII

hpy188I

mbolI

bpuAI

hpaII

bsi

hpaII

hpy188I

mbolI

bpuAI

hpaII

bsi

hpaII

hpy188I

mbolI

bpuAI

hpaII

bsi

hpaII

hpy188I

mbolI

bpuAI

hpaII

bsi

hpaII

hpy188I

mbolI

bpuAI

hpaII

bsi

hpaII

hpy188I

mbolI

bpuAI

hpaII

bsi

hpaII

hpy188I

mbolI

bpuAI

hpaII

bsi

hpaII

hpy188I

mbolI

bpuAI

hpaII

bsi

hpaII

hpy188I

mbolI

bpuAI

hpaII

bsi

hpaII

hpy188I

mbolI

bpuAI

hpaII

bsi

hpaII

hpy188I

mbolI

bpuAI

hpaII

bsi

hpaII

hpy188I

mbolI

bpuAI

hpaII

bsi

hpa

tth1111/aspi

plei

pf1f1

mly1

hinf1

bsma1 bsma1

bseR1

mul1

bseR1

bpm1/gsu1[dcn-]

hinf1 dde1

hha1/efo1 bspcni

bst4cl/hpych4111 mul1 hpych4v

1601 AGCAAGACAG TTGTTTCTCC TCCTCCTTGC ATATTCTTA CTCGCTCCA GCCGAGTGA TAGAGTGA CTCGTCTCA AAAAAAGTA TCTTAATA

TCGTTCTGTC AACAAAGAG AGAGGAAGC TATAAGCAT GAGCGAGGT CGGACTCACT ATCTCACTCT GAGACAGAGT TTTTTCAT AGAGATTAT

tru91

mse1

tsp5091

ps11

sml1

1701 CAGATTATA ATTCTGCTT GAGTATGTG TTAAGTACT TGTATTAGA AGATTTCAG ATTCAATCCA TCCTCTTAGT TTTCTTTTAA GTGACCCAT

GTCTTAATAT TAAAGACGAA CTCATACCAC AATTGATGA ACATAATCT TTCTAAAGTC TAAGTAAAGT AGAGGAATCA AAGAAAATT CCACTGGGA

dde1[M.alu1-]

alu1

1801 CTGTGATAAA AATATAGCTT AGTGTAAAA TCAGTGTAC TTATACATGG CCTAAATGT TTCTACAAAT TAGAGTTGT CACTTATTC ATTGTACCT

GACACTATTT TTATATCGAA TCACGATTTT AGTCACATG AATATGTACC GGATTTTACA AAGATGTTA ATCTCAACA GTGAATAAG TAAACATGA

mae111

tspR1

alu1

1801 CTGTGATAAA AATATAGCTT AGTGTAAAA TCAGTGTAC TTATACATGG CCTAAATGT TTCTACAAAT TAGAGTTGT CACTTATTC ATTGTACCT

GACACTATTT TTATATCGAA TCACGATTTT AGTCACATG AATATGTACC GGATTTTACA AAGATGTTA ATCTCAACA GTGAATAAG TAAACATGA

tsp451

tsp5091

mae111

csp61

rsal

hph1

tru91 mae111

mse1 bste11

dde1

asp700

hpy1881

hinf1

xm1

hinc11/hind11

hpa1

trp91

tsp451

mae111

hph1

trp91

tsp451

mae111

hph1

trp91

tsp451

mae111

hph1

trp91

```

scrFI[dcM-]
pspGI
mvaI
ecorII[dcM-]
dsav[dcM-]
bstNI
haeIII/paII
msci/baII[dcM-]
eaeI[dcM-]
cfrI
scrFI[dcM-]
pspGI
mvaI bskI[dcM-]
ecorII[dcM-] tsp45I
dsav[dcM-] maeIII
bstNI hinfI
bskI[dcM-] tspRI
pleI bslI[dcM-] hhaI/cfoI
mlyI bsaJI apyI[dcM+]
hinfI apyI[dcM+] btsI
ddeI
bspcNI
1901 AAGAGAAAA TAGGCTCAGT TAGAAAGA CTCCTGGCC AGCGCAGTG ACTTACGCTT GTAATCTCAG CACTTGGGA GGCCAAGCA GGCAGATCAC
TTCTCTTTT ATCCGAGTCA ATCTTTCCCT GAGGACCGG TCCCGCTCAC TGAATGCGGA CATTAGAGTC GTGAACCCCT CCGTTCGT CCGTCTAGTG

```

msci/bali[dcn-]  
eaei[dcn-]

scrfl[dcn-]

pspgi

mval

ecoriI[dcn-]

dsav[dcn-]

bstNI

bsmAI bsski[dcn-]

taqI foki cfrI nlaIII

bsmAI

hpy188III bsaI bsrFSI haeIII/palI

esp3I

mnlI hpy188III apyI[dcn+] hphI

bsmBI

tsp509I

alul

nlaIV

2001 GAGGTACAGA GTTCGAGACC ATCCTGGCCA ACATGTGTA ACCCGCTCTC TACTAAAT ATAAATTA GCTGGGTGTG GTGGCAGAG CCTGTATCC  
CTCAGTCTCT CAAGCTCTGG TAGGACCGGT TGTACACTT TGGGCAGAG ATGATTTTAA TATTTTAAAT CGACCCACAC CACCGTCTC GGACATTAGG

scrfl[dcn-]

pspgi

mval

ecoriI[dcn-]

dsav[dcn-]

bstNI

bski[dcn-]

apyl[dcn+]

apyl[dcn+]

2101 CAGCTACACA GGAGGCTGAG GCACAGAAAT CACTTGAAT CAGGAGATGG AGTTTCACT GAGCCGAGAT CAGCCACTG CACTCCAGCC TGGCAACAGA  
GTCGATGTGT CCTCCACTC CGTCTCTTA GTGAAGTGA GTCCCTACAC TCAGAGTCA CTCGGCTCTA GTGGGTGAC GTGAGTGG ACCGTGTCT

ddel tfil

hpy188III

bspcNI hinfI

ddel

alul mnlI mnlI bssSI

bspcNI

mnlI tspRI

dpnl[dan+] bsgI bpnI/gsuI[dcn-]

2101 CAGCTACACA GGAGGCTGAG GCACAGAAAT CACTTGAAT CAGGAGATGG AGTTTCACT GAGCCGAGAT CAGCCACTG CACTCCAGCC TGGCAACAGA  
GTCGATGTGT CCTCCACTC CGTCTCTTA GTGAAGTGA GTCCCTACAC TCAGAGTCA CTCGGCTCTA GTGGGTGAC GTGAGTGG ACCGTGTCT

```

fnu4HI/bsoFI
haeIII/palI
mcrI
eagI/xmaIII/ecI XI
eaeI
cfrI
bsrI rmaI
notI maeI
fnu4HI/bsoFI bfaI
acI acI speI
bsmAI
2201 GCGAGACTCC ATCTCAAAA AAAAAAAAAA AAAAAAAAAA AAAAAAAAAAGG CGGCCGCCGA CTAGTGAGC
CGCTCTGAGG TAGAGTTTT TTTTTTTTT TTTTTTTTT TTTTTTTCC GCCGCGGCT GATCACTCG

```

> length: 2269

```

accI (GTMKAC) : 1501
acI (CCGC) : 39 498 2250 2254
afI (ACRYGT) : 780 1586
ahaIII (TTTAA) : 1150
ahdI (GACNNNNNGTC) : 278 714
aluI (AGCT) : 152 300 429 510 690 822 888 1015 1345 1476 1816 2070 2102
alw26I (CAGNNNCTG) : 101 316
alwI (GATCNNNN) : 318 530
alwNI (CAGNNNCTG) : 101 316
apoI (RAATY) : 3 310 423 655 1464
apyI (CCWGG) : 321 332 1422 1934 1939 2023 2189
asp700 (GAANNNTTC) : 1464 1749
asphI (GWCWC) : 1582

```

GENE GENES										SYSTEMS		Additional Resource:			
GENE VIEWER		GENE		FAM		MAP		GENE HUB		Find		New		Update	
SEQUENCE VIEWER		DNA		GRC		FNA		IS		FLS		OLI		SELECT	
ASSAY VIEWER		PROC		DOM		EXP		PUR		LOT		ASY		GO	

## DNA64883

DNA Info [Project DNA56011](#)Is Primarydna ☒Source Info [628 FLS 2333](#) [LIB309](#) [RNA217](#) [SRC685](#) Human blank IncyteGene Info [PRO1244](#) Human [IAG2](#) [UNQ628](#)

## Gene Annotation

Genome Mapping [Run Geode](#)

Affy [HGU133A 221553\\_at](#)  
[HGU133B 224899\\_s\\_at](#)  
[HGU133P 221553\\_at, 224899\\_s\\_at](#)  
[HGU95B 43481\\_at, 45230\\_at](#)  
[HGU95E 90925\\_at](#)  
[Hu35KB RC N48787\\_at](#)  
[Hu35KC RC\\_AA133886\\_at, RC\\_AA157267\\_at](#)  
[HuGenen1 AF130104\\_at, P\\_A37058\\_at](#)  
[MGU74B 106968\\_at, 110460\\_at](#)  
[MGU74C 140441\\_at](#)  
[MOE430A 1419459\\_a\\_at](#)  
[MOE430P 1419459\\_a\\_at](#)

Agilent [H1Av2 A\\_23\\_P148273](#)  
[H1A A\\_23\\_P148273](#)  
[H1Av2 A\\_23\\_P148273](#)  
[M1A A\\_51\\_P206624](#)  
[WHG A\\_23\\_P148273, A\\_24\\_P105164, A\\_24\\_P333802, A\\_24\\_P485219](#)

FANTOM [Mouse:2410001C15, 2610529C04, 2810435A10, 9130015A07](#)GenBank [Human:AF130104, AK027632, AK075394, AL136636, AY358691, BC060842, BC063037](#)  
[Mouse:AK010320, AK012185, AK013243, AK018623, BC003881](#)GeneHub [Human:GENE30707](#)  
[Mouse:MGENE40317](#)INCYFL [Human:959272.FL1\\_0](#)Incyte [Human:1397153.5, 1397153.6](#)LocusLink [Human:84061](#)  
[Mouse:67075](#)MGI [Mouse:1914325](#)Proteome [Human:NP\\_115497.3](#)  
[Mouse:NP\\_080228.2](#)



UniGene Human:Hs.323562  
Mouse:Mm.275943

DNA64883 [REDACTED]  
DNA64883 [REDACTED]  
DNA64883 [REDACTED]  
DNA64883 [REDACTED]  
DNA64883 [REDACTED]  
DNA64883 [REDACTED]  
DNA64883 [REDACTED]  
DNA64883 [REDACTED]  
DNA64883 [REDACTED]

DNA64883 [REDACTED]  
DNA64883 [REDACTED]  
DNA64883 [REDACTED]  
DNA64883 [REDACTED]  
DNA64883 [REDACTED]

**Status**Scientist Alane Gray

Notebook 28923

Page 86

Storage Location

Box

Slot

Inventory Status

Others ☐ Sent to pLASMID Archive  
☐ Clone Verified

Date Entered [REDACTED]

Date Updated [REDACTED]

Date Completed

Date Canceled

Cancel Reason

Clone Status not reviewed

Sequence Status

**Project Member**

No Project member generated

**FLS FLSDNA**

No FLS, FLSDNA generated

**Exp Construct**

EXP	Lab Name	Construct DNA	System
<u>EXP2003</u>	PH.64883.his.163	<u>DNA75635</u>	Baculovirus
<u>EXP1945</u>	sst.64883.H8	<u>DNA79249</u>	Mammalian Stable
<u>EXP2354</u>		<u>DNA79249</u>	Mammalian Stable
<u>EXP2342</u>	64883.644 IgG	<u>DNA83540</u>	Baculovirus

**ABI**

No ABI run generated

**MA Plate**

MA Plate	Well Num	Well Location	Date	Typ Plate
PLT72	61	F1	[REDACTED]	Storage

**Print Run**

No Print run generated

**XPT**

No XPT generated

[ASY](#) | [DNA](#) | [DOM](#) | [EXP](#) | [FAM](#) | [FLS](#) | [LIB](#) | [LOT](#) | [MAP](#) | [OLI](#) | [PRB](#) | [PRO](#) | [PUR](#) | [RNA](#) | [SRC](#) | [UNQ](#) | [XPT](#) | [YST](#)  
[Assay Viewer](#) | [Sequence Viewer](#) | [Gene Viewer](#) | [GenenGenes](#) | [SAGE](#)

GenenGenes Feedback

**This Page is Inserted by IFW Indexing and Scanning  
Operations and is not part of the Official Record**

**BEST AVAILABLE IMAGES**

Defective images within this document are accurate representations of the original documents submitted by the applicant.

Defects in the images include but are not limited to the items checked:

- ☐ **BLACK BORDERS**
- ☐ **IMAGE CUT OFF AT TOP, BOTTOM OR SIDES**
- ☐ **FADED TEXT OR DRAWING**
- ☐ **BLURRED OR ILLEGIBLE TEXT OR DRAWING**
- ☐ **SKEWED/SLANTED IMAGES**
- ☐ **COLOR OR BLACK AND WHITE PHOTOGRAPHS**
- ☐ **GRAY SCALE DOCUMENTS**
- ☐ **LINES OR MARKS ON ORIGINAL DOCUMENT**
- ☐ **REFERENCE(S) OR EXHIBIT(S) SUBMITTED ARE POOR QUALITY**
- ☐ **OTHER:** \_\_\_\_\_

**IMAGES ARE BEST AVAILABLE COPY.**

**As rescanning these documents will not correct the image problems checked, please do not report these problems to the IFW Image Problem Mailbox.**